

IN THE CLAIMS

Please amend the claims as follows:

1. (Cancelled).
2. **(Currently Amended)** A method of presenting an antigenic peptide on the surface of a viable cancer cell, said method comprising:
 - contacting said cancer cell with said antigenic peptide and with a photosensitizing agent *ex vivo*, wherein said peptide and said agent are each taken up into an intracellular membrane-restricted compartment of said cell;
 - irradiating said cell *ex vivo* with light of a wavelength effective to activate the photosensitizing agent, such that the membrane of said intracellular compartment is disrupted, releasing said peptide into the cytosol of the cell, without killing the cell;
 - wherein, said released antigenic peptide, or a part thereof of sufficient size to stimulate a cytotoxic T cell response, is subsequently presented on the surface of said cell by a class I MHC molecule;
 - administering the cell to a mammal after irradiating said cell;
 - wherein presentation of the antigenic peptide, or part thereof, on the surface of said cell results in cytotoxic T cell mediated cell killing by a cytotoxic T cell specific for said antigenic peptide or a part thereof; and
 - wherein the photosensitizing agent is selected from the group consisting of a porphyrin, phthalocyanine and a chlorin.
3. (Cancelled).
4. (Previously Presented) The method of claim 2, wherein the antigenic peptide is a vaccine antigen or vaccine component.
- 5-7. (Cancelled).

8. (Previously Presented) The method of claim 2 wherein the photosensitizing agent is meso-tetraphenylporphine with 4 sulfonate groups (TPPS₄), meso-tetraphenylporphine with 2 sulfonate groups on adjacent phenyl rings (TPPS_{2a}), or aluminum phthalocyanine with 2 sulfonate groups on adjacent phenyl rings (AlPcS_{2a}).
9. (Previously Presented) The method of claim 2, wherein the antigenic peptide and/or photosensitizing agent is bound to one or more targeting agents or carrier molecules.
10. (Canceled) .
- 11-27. (Canceled).
28. (Previously Presented) The method of claim 2, wherein at least 90% of the cells are not killed.
29. (Previously Presented) The method of claim 2, wherein at least 95% of the cells are not killed.
30. (Previously Presented) The method of claim 2, wherein the photosensitizing agent is a sulfonated tetraphenylporphine, a disulfonated aluminum phthalocyanine or a tetrasulfonated aluminum phthalocyanine.
31. (Canceled)
32. (Canceled)
- 33-36. (Canceled)

37. (Cancelled)

38-40. (Canceled)

41. (Previously Presented) The method of claim 2, wherein the antigenic peptide stimulates cytotoxic T cells.

42. (Canceled).

43. (New) An *in vitro* method of presenting an antigenic peptide on the surface of a viable cancer cell and killing said cell by cytotoxic T cell mediated cell killing, said method comprising:

contacting said cancer cell with said antigenic peptide and with a photosensitizing agent, wherein said peptide and said agent are each taken up into an intracellular membrane-restricted compartment of said cell;

irradiating said cell with light of a wavelength effective to activate the photosensitizing agent, such that the membrane of said intracellular compartment is disrupted, releasing said peptide into the cytosol of the cell, without killing the cell;

wherein said released antigenic peptide, or a part thereof of sufficient size to stimulate a cytotoxic T cell response, is subsequently presented on the surface of said cell by a class I MHC molecule;

wherein presentation of the antigenic peptide, or part thereof, on the surface of said cell results in cytotoxic T cell mediated cell killing by a cytotoxic T cell specific for said antigenic peptide or a part thereof; and

wherein the photosensitizing agent is selected from the group consisting of a porphyrin, phthalocyanine and a chlorin.

44. (New) The method of claim 43, wherein the antigenic peptide is a vaccine antigen or vaccine component.

45. (New) The method of claim 43, wherein the photosensitizing agent is meso-tetraphenylporphine with 4 sulfonate groups (TPPS₄), meso-tetraphenylporphine with 2 sulfonate groups on adjacent phenyl rings (TPPS_{2a}), or aluminum phthalocyanine with 2 sulfonate groups on adjacent phenyl rings (AlPcS_{2a}).
46. (New) The method of claim 43, wherein the antigenic peptide and/or photosensitizing agent is bound to one or more targeting agents or carrier molecules.
47. (New) The method of claim 43, wherein at least 90% of the cells are not killed.
48. (New) The method of claim 43, wherein at least 95% of the cells are not killed.
49. (New) The method of claim 43, wherein the photosensitizing agent is a sulfonated tetraphenylporphine, a disulfonated aluminum phthalocyanine or a tetrasulfonated aluminum phthalocyanine.
50. (New) The method of claim 43, wherein the antigenic peptide stimulates cytotoxic T cells.